

Please amend claim8 as follows:

- 1           8. (Amended) Pharmaceutical composition according to claim 1 further comprising at least
- 2 one corticosteroid.

Please amend claim 12 as follows:

- 1           12. (Amended) Pharmaceutical composition according to claim 1, wherein adjuvants and
- 2 excipients are ethanol and sodium chloride.

Please amend claim 13 as follows:

- 1           13. (Amended) Pharmaceutical composition according to claim 1 for use in the treatment
- 2 of cutaneous injuries.

Please amend claim 14 as follows:

- 1           14. (Amended) Pharmaceutical composition according to claim 1 for use in the treatment
- 2 of cutaneous injuries resulting from mechanical traumas or surgical operations, burns, dermatitis
- 3 both from animal sting and animal or poisonous plant contact.

Please amend claim 15 as follows:

- 1           15. (Amended) Pharmaceutical composition according to claim 1 for use in the therapy
- 2 and prevention of hypertrophic cicatrices and keloids.

Please amend claim16 as follows:

- 1           16. (Amended) Pharmaceutical composition according to claim 1 for use in aesthetic
- 2 medicine.

Please amend claim 18 as follows:

- 1           18. (Amended) Pharmaceutical composition according to claim 1 in the form selected  
2 from the group consisting of ointment, gel, foam, liquid preparations, medicated plaster.

Please amend claim 19 as follows:

- 1           19. (Amended) A process for the preparation of a pharmaceutical composition comprising  
2 the following steps: a) preparing in an anhydrous atmosphere, distinct A and B mixtures,  
3 respectively, of (A) trichloroacetic acid at a concentration from about 40 % to about 90 % in a  
4 suitable polymer and (B) 2-hydroxybenzoic acid at a concentration from about 20 % to about 60  
5 %, in a suitable polymer; b) mixing by adding small subsequent portions, same volumes of the two  
6 mixtures to obtain the A+B mixture; c) adding a volume of a (1 $\alpha$ , 2 $\beta$ , 5 $\alpha$ )-5-methyl-2-(1-  
7 methylethyl)cyclohexanol saturated solution in anhydrous ethanol equal to 2 % of the volume of  
8 the A+B mixture; d) adding sodium chloride in a such amount to obtain a final concentration of  
9 about 1,2 % w/v; e) placing the obtained composition in a bottle filled with anhydrous air at 30°C  
10 in reduced pressure conditions and at ambient temperature for about 24 hours.

Please amend claim 22 as follows:

22. (Amended) Process according to claim 19, wherein to the mixture obtained in step d)  
one or more corticosteroids is added.